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IN THE CLAIMS:

Kindly amend the claims, without prejudice, without admission, without surrender of subject matter, and without any intention of creating any estoppel as to equivalents, to read as follows:

- 1. (Currently Amended) A composition comprising a coagulation factor IXa FIXa and a composition comprising a coagulation factor VIII FVIII for simultaneous, simultaneous separate or sequential use in the treatment of haemophilia A or haemophilia B in a subject which who does not present with anti-FVIII antibodies.
- 2. (Currently Amended) A method of using a coagulation factor IXa and a coagulation factor VIII FIXa and FVIII in the preparation of a composition according to claim 1 for the treatment of haemophilia A or haemophilia B in a subject which does not present with anti-coagulation factor VIII anti-FVIII anti-bodies wherein the method comprises admixing coagulation factors IXa and VIII.
- 3. (Original) A composition comprising FIXa according to claim 1, which further comprises phospholipid.
- 4. (Original) A method of using FIXa in the manufacture of a composition comprising FVIII for the treatment of haemophilia A or haemophilia B, wherein the presence of FIXa allows the concentration of FVIII in the composition to be reduced in comparison to a composition which does not comprise FIXa.
- 5. (Currently Amended) The method according to claim 4, wherein the composition is administered to a subject which who does not present with anti-FVIII antibodies.
- 6. (Original) The method according to claim 4, wherein the FVIII and FIXa reagents are produced using recombinant DNA technology.

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- 7. (Original) The method according to claim 4, wherein the composition further comprises phospholipid.
- 8. (Original) The method according to claim 4, wherein the composition is formulated to provide FVIII to a subject at a dosage of between 2 and 10 IU/kg.
- 9. (Withdrawn) A method for treating a subject suffering from haemophilia A or haemophilia B, comprising administering to a subject in need thereof a composition comprising FIXa and FVIII, wherein said subject does not present with anti-FVIII antibodies or wherein said composition comprises FVIII in an amount lower than that required for treatment of said subject with a composition lacking FIXa.
- 10. (Withdrawn) The method according to claim 9, wherein said composition further comprises phospholipid.
- 11. (Withdrawn) The method according to claim 9, wherein the composition comprises recombinant FIXa and recombinant FVIII.
- 12. (Withdrawn) The method according to claim 9, wherein the composition is formulated to provide FVIII to a subject at a dosage of between 2 and 10 IU/kg.
- 13. (Original) A method for potentiating FVIII comprising the step of mixing together Factor FVIII and FIXa into a composition.
- 14. (Original) The method according to claim 13, wherein said composition further comprises phospholipid.
- 15. (Original) The method according to claim 13, wherein the composition comprises recombinant FIXa and recombinant FVIII.

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- 16. (Withdrawn) A method for reducing the immunogenicity of FVIII in a composition comprising FVIII in a subject, comprising administering FVIII together with FIXa to the subject.
- 17. (Withdrawn) A method of using FIXa and FVIII in the preparation of a composition for the treatment of haemophilia, wherein the FVIII in said composition has reduced immunogenicity as a result of the presence of FIXa.